

**K960475 MODIFIED LUXAR LX-20 FAMILY CO2 LASER SYSTEM**Apr 26, 1996  
85 days to decisionK960475 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k960475/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Feb 1, 1996
Decision date	Apr 26, 1996
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Luxar Corp.</b>
Location	Bothell, WA, US
Contact	JONATHAN S KAHN
510(k) history	17 submissions · 17 cleared · 1988-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960475/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 5, 2026